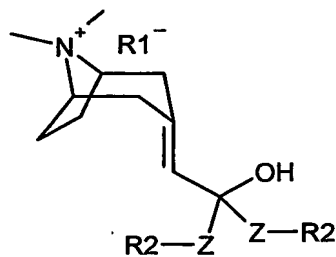
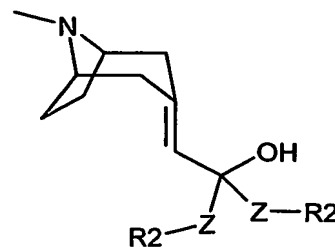


What is claimed is:

1. A compound having structure I or II as indicated below:



(I)



(II)

- 5 wherein:

R1<sup>-</sup> represents an anion associated with the positive charge of the N atom;

- R2 is selected from the group consisting of cycloalkyl groups having from 5 to 6 carbon atoms, cycloalkyl-alkyl having 6 to 10 carbon atoms, heterocycloalkyl having 5 to 6 carbon atoms and N or O as the heteroatom, heterocycloalkyl-alkyl having 6 to 10 carbon atoms and N or O as the heteroatom, aryl, optionally substituted aryl, heteroaryl, and optionally substituted heteroaryl; and  
 10 Z is a bond or (C<sub>1</sub>-C<sub>6</sub>)alkyl.

2. A compound according to claim 1 wherein R1<sup>-</sup> is selected from the group consisting of chloride, bromide, iodide, sulfate, benzene sulfonate and toluene sulfonate.  
 15

3. A compound according to claim 1 selected from the group consisting of:  
 2-(8-Methyl-8-aza-bicyclo[3.2.1]oct-3-ylidene)-1,1-di-thiophen-2-yl-ethanol;  
 2-Benzyl-1-(8-methyl-8-aza-bicyclo[3.2.1]oct-3-ylidene)-3-phenyl-propan-2-ol;  
 20 2-(8-Methyl-8-aza-bicyclo[3.2.1]oct-3-ylidene)-1,1-diphenyl-ethanol;  
 3-(2-Hydroxy-2,2-di-thiophen-2-yl-ethylidene)-8,8-dimethyl-8-azonia-bicyclo[3.2.1]octane iodide;  
 3-(2-Benzyl-2-hydroxy-3-phenyl-propylidene)-8,8-dimethyl-8-azonia-bicyclo[3.2.1]octane iodide;  
 25 3-(2-Hydroxy-2,2-diphenyl-ethylidene)-8,8-dimethyl-8-azonia-bicyclo[3.2.1]octane iodide.

4. A compound according to claim 3 selected from the group consisting of:  
3-(2-Hydroxy-2,2-di-thiophen-2-yl-ethylidene)-8,8-dimethyl-8-azonia-  
bicyclo[3.2.1]octane iodide;  
3-(2-Benzyl-2-hydroxy-3-phenyl-propylidene)-8,8-dimethyl-8-azonia-  
5 bicyclo[3.2.1]octane iodide; and  
3-(2-Hydroxy-2,2-diphenyl-ethylidene)-8,8-dimethyl-8-azonia-bicyclo[3.2.1]octane  
iodide.
5. A pharmaceutical composition for the treatment of muscarinic acetylcholine  
10 receptor mediated diseases comprising a compound according to claim 1 and a  
pharmaceutically acceptable carrier thereof.
6. A method of inhibiting the binding of acetylcholine to its receptors in a  
mammal in need thereof comprising administering a safe and effective amount of a  
15 compound according to claim 1.
7. A method of treating a muscarinic acetylcholine receptor mediated disease,  
wherein acetylcholine binds to said receptor, comprising administering a safe and  
effective amount of a compound according to claim 1.  
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8. A method according to claim 7 wherein the disease is selected from the group  
consisting of chronic obstructive lung disease, chronic bronchitis, asthma, chronic  
respiratory obstruction, pulmonary fibrosis, pulmonary emphysema and allergic  
rhinitis.  
25
9. A method according to claim 7 wherein administration is via inhalation via the  
mouth or nose.
10. A method according to claim 7 wherein administration is via a medicament  
30 dispenser selected from a reservoir dry powder inhaler, a multi-dose dry powder inhaler  
or a metered dose inhaler.

11. A method according to claim 8 wherein the compound is administered to a human and has a duration of action of 12 hours or more for a 1 mg dose.
12. A method according to claim 11 wherein the compound has a duration of action  
5 of 24 hours or more.
13. A method according to claim 12 wherein the compound has a duration of action of 36 hours or more.